The isolated HLA DRB1*15-binding peptide of claim 2 wherein the

5.(twice amended)

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isolated peptide comprises an endosomal targeting signal. The isolated HLA DRB1*15-binding peptide of claim 2 wherein the

7.(twice amended)

isolated peptide is non-hydrolyzable.

A composition comprising an isolated MAGE-Al HLA class I-

binding peptide and an isolated MAGE-A1 HLA DRB1*15-binding peptide, wherein the 9.(twice amended)

isolated HLA class II-binding peptide consists of

the amino acid sequence set forth as SEQ ID NO:7, or a functional variant thereof 0-10 amino acids added to either or both ends of the amino acid sequence set forth as consisting of one amino acid addition, substitution or deletion, and

SEQ ID NO:7, or the functional variant thereof consisting of one amino acid addition,

14.(amended) The composition of claim 9 wherein the isolated MAGE-Al HLA DRB1*15substitution or deletion.

binding peptide comprises an endosomal targeting signal.

The isolated HLA DRB1*15-binding peptide of claim 2 wherein the

isolated peptide consists of an amino acid sequence selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4, and SEQ ID NO:7, or a functional variant thereof consisting of one 76.(twice amended)

77.(amended) The isolated HLA DRB1*15-binding peptide of claim 2 wherein the isolated amino acid addition, substitution or deletion.

peptide consists of an amino acid sequence selected from the group consisting of SEQ ID NO:3,

78.(amended) The isolated HLA DRB1*15-binding peptide of claim 5 wherein the endosomal SEQ ID NO:4 and SEQ ID NO:7.

targeting signal comprises an endosomal targeting portion of human invariant chain Ii or LAMP-

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79.(amended) The isolated HLA DRB1*15-binding peptide of claim 7 wherein the isolated peptide is selected from the group consisting of peptides comprising D-amino acids, peptides comprising a -psi[CH₂NH]-reduced amide peptide bond, peptides comprising a -psi[COCH₂]-ketomethylene peptide bond, peptides comprising a -psi[CH(CN)NH]-(cyanomethylene)amino peptide bond, peptides comprising a -psi[CH₂CH(OH)]-hydroxyethylene peptide bond, peptides comprising a -psi[CH₂O]-peptide bond, and peptides comprising a -psi[CH₂S]-thiomethylene peptide bond.

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80.(amended) The composition of claim 9 wherein the MAGE-A1 HLA class I-binding peptide and the MAGE-A1 HLA DRB1*15-binding peptide are combined as a polytope polypeptide.

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81.(twice amended) The composition of claim 9 wherein the isolated MAGE-A1 HLA DRB1*15-binding peptide consists of an amino acid sequence selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4 and SEQ ID NO:7, or a functional variant thereof consisting of one amino acid addition, substitution or deletion.

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82.(twice amended) The composition of claim 9 wherein the isolated MAGE-A1 HLA DRB1*15-binding peptide consists of an amino acid sequence selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4 and SEQ ID NO:7.

<u>Remarks</u>

Applicants have amended claims 2, and 9 to clarify the subject matter of the claims with respect to the scope of the functional variants and the nature of the peptides as HLA DRB1*15-binding peptides. The remaining claims were amended to reflect the amendments to claims 2 and 9. No new matter has been added.

Regarding the Examiner's statement in paragraph 4 of the Office Action, Applicants respectfully disagree that the transitional phrase "consisting essentially of" is an open term. To the contrary, it is well settled in case law that "consisting essentially of" has a meaning distinct from open terms such as "comprising." As interpreted by the courts, consisting essentially of covers the claimed article with additional elements, but excludes additional elements "that would materially affect the basic and novel characteristics" of the products defined in the balance of